

Fate of *Salmonella* Typhimurium in laboratory-scale drinking water biofilms

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Abstract

Investigations were carried out to evaluate and quantify colonization of laboratory-scale drinking water biofilms by a chromosomally green fluorescent protein (gfp)-tagged strain of *Salmonella* Typhimurium. Gfp encodes the green fluorescent protein and thus allows in situ detection of undisturbed cells and is ideally suited for monitoring *Salmonella* in biofilms. The fate and persistence of non-typhoidal *Salmonella* in simulated drinking water biofilms was investigated. The ability of *Salmonella* to form biofilms in monoculture and the fate and persistence of *Salmonella* in a mixed aquatic biofilm was examined. In monoculture *S. Typhimurium* formed loosely structured biofilms. *Salmonella* colonized established multi-species drinking water biofilms within 24 hours, forming micro-colonies within the biofilm. *S. Typhimurium* was also released at high levels from the drinking water-associated biofilm into the water passing through the system. This indicated that *Salmonella* could enter into, survive and grow within, and be released from a drinking water biofilm. The ability of *Salmonella* to survive and persist in a drinking water biofilm, and be released at high levels into the flow for recolonization elsewhere, indicates the potential for a persistent health risk to consumers once a network becomes contaminated with this bacterium.